

Flourescence Signal

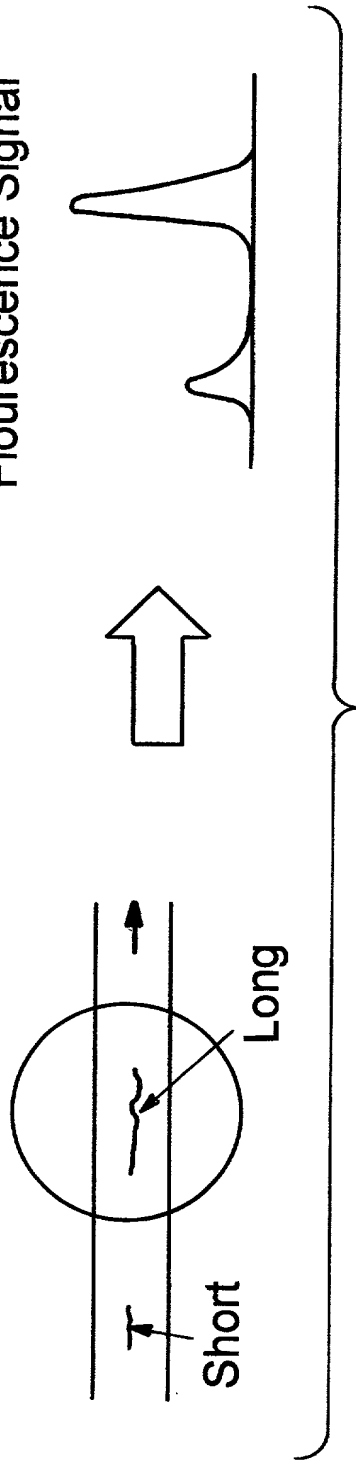


Fig. 1A

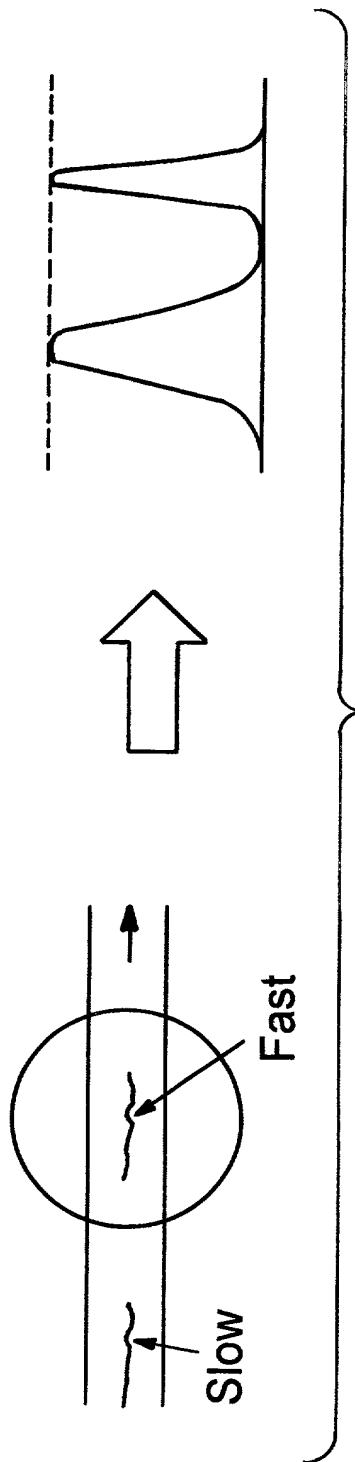


Fig. 1B

VIM - system

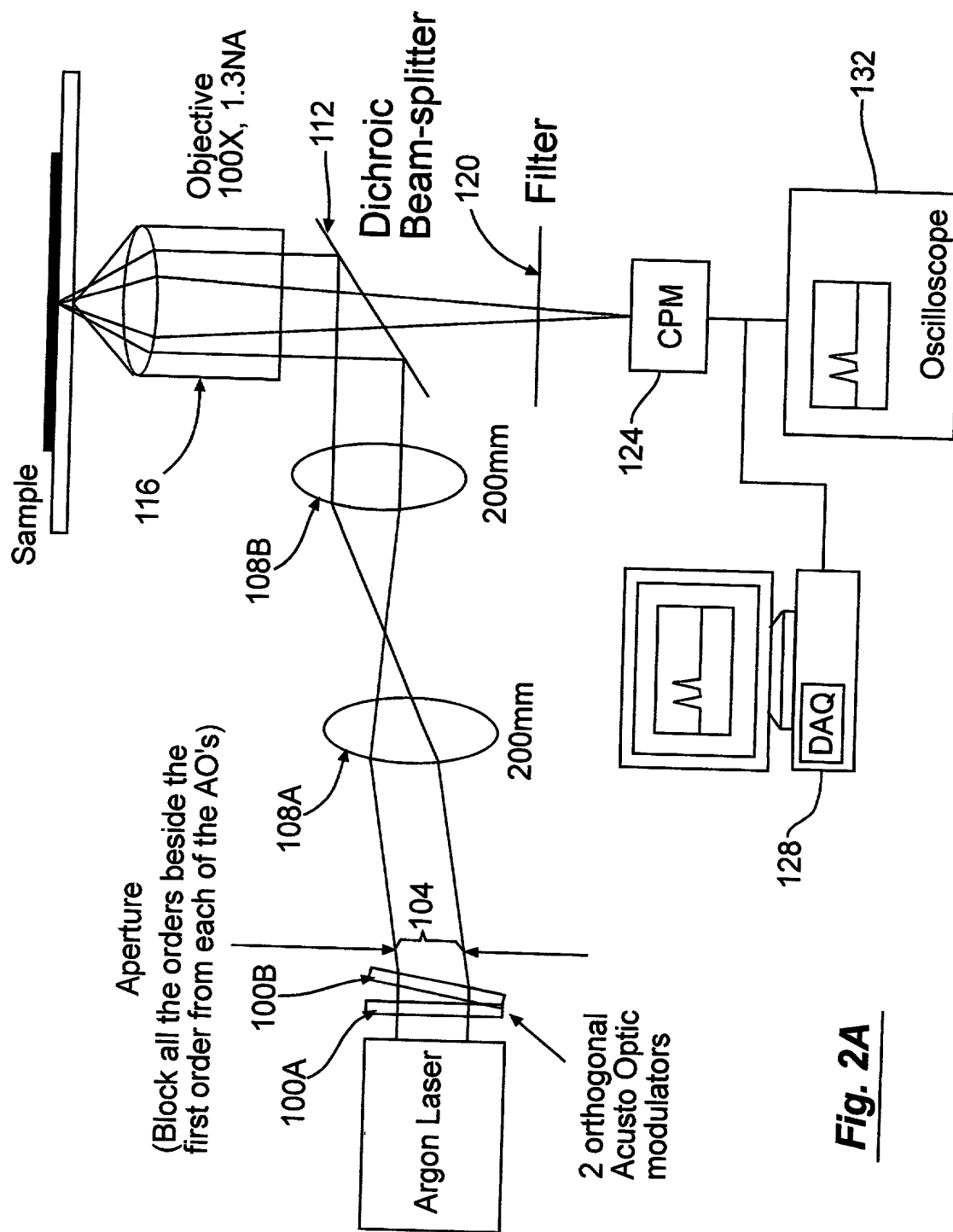


Fig. 2A

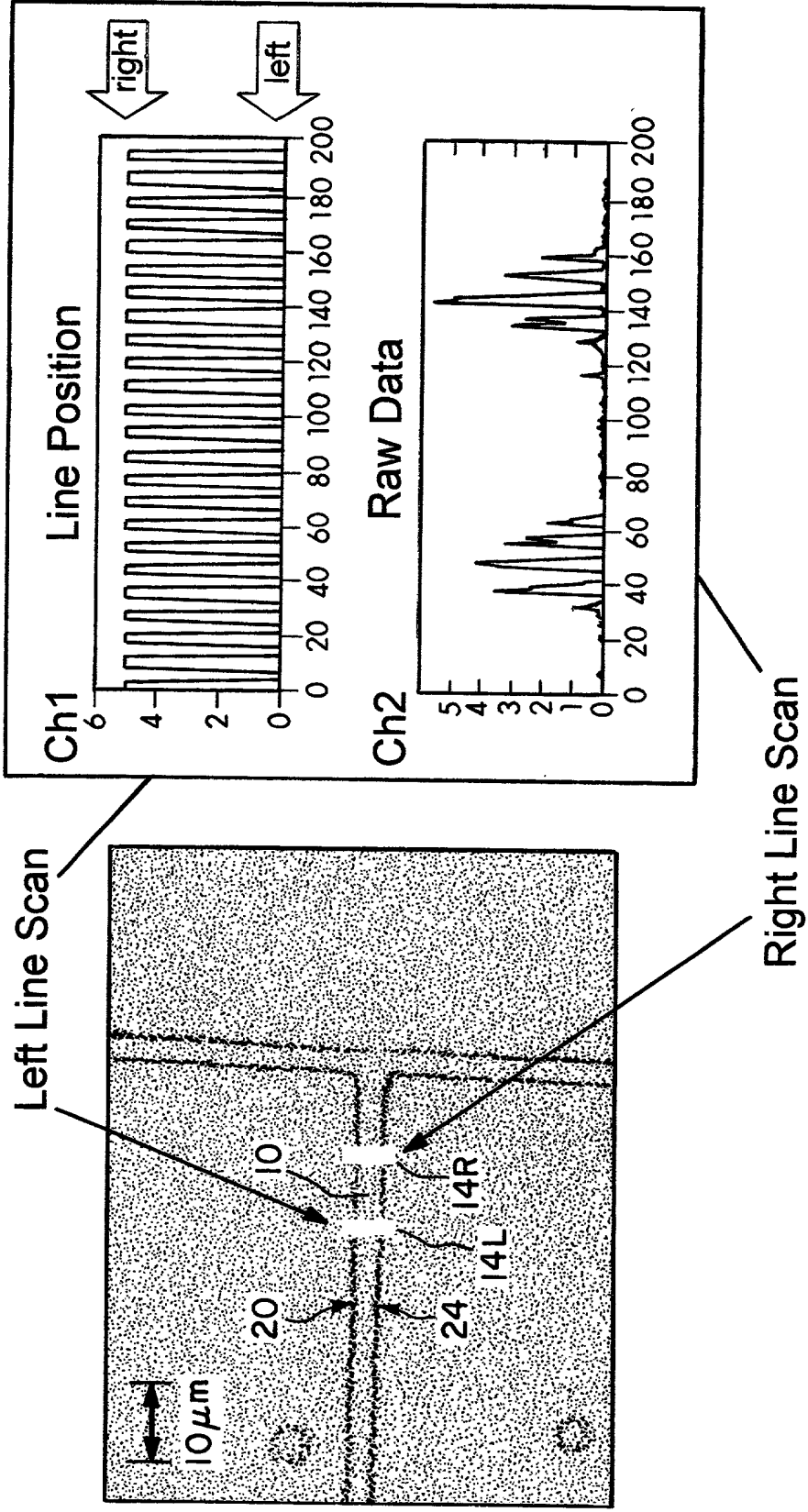


Fig. 2B

The beam after the two Acusto Optics Modulators

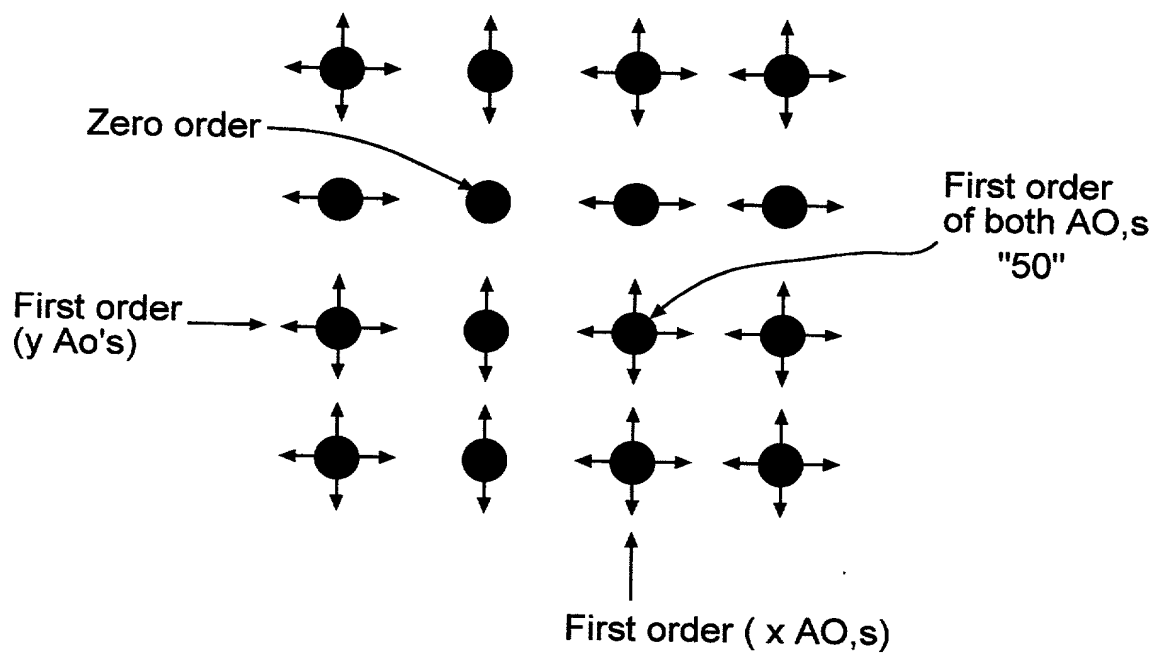


Fig. 2C

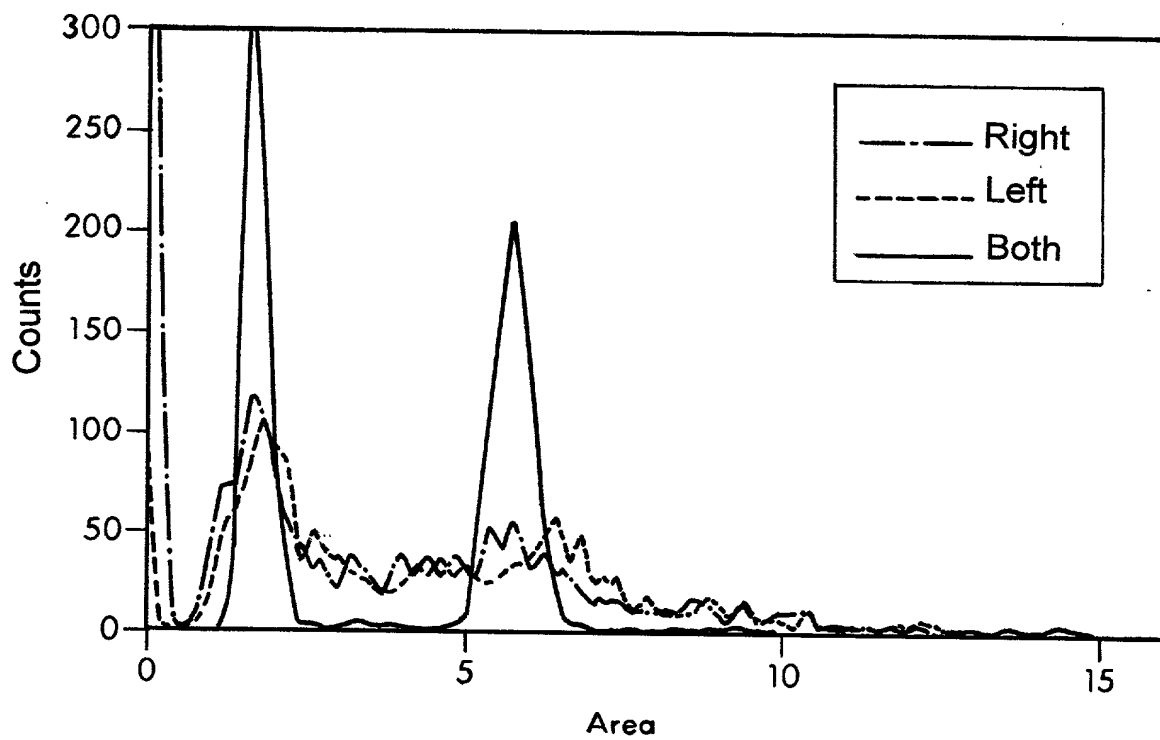


Fig. 3

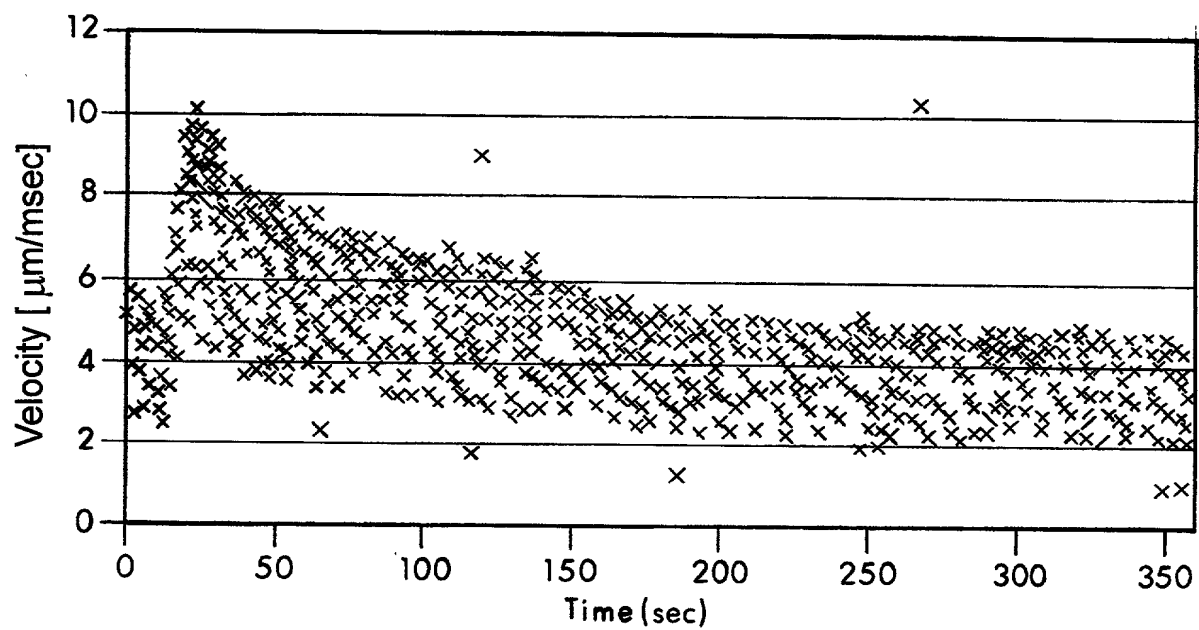


Fig. 4

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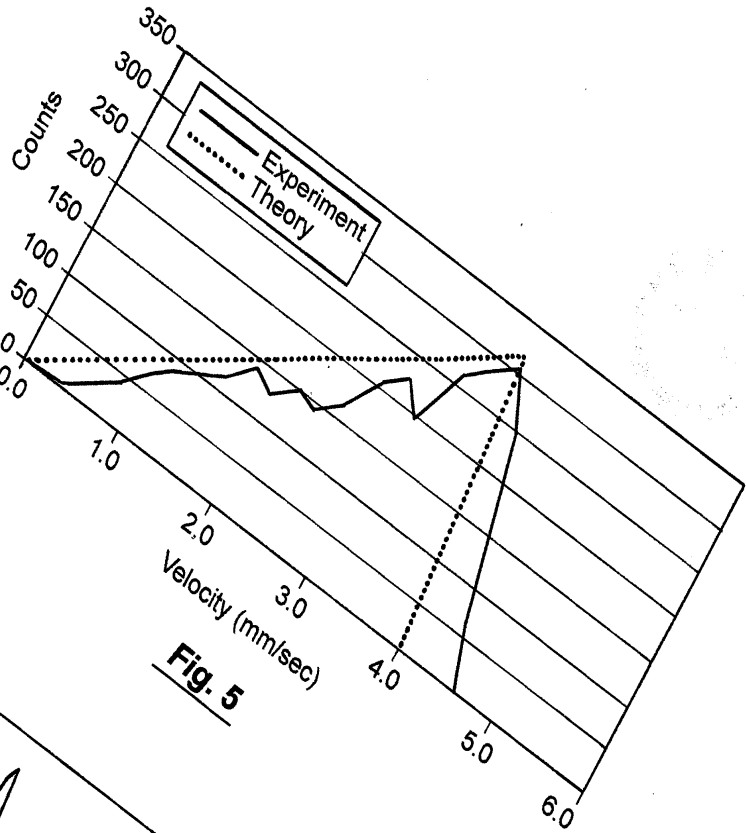


Fig. 5

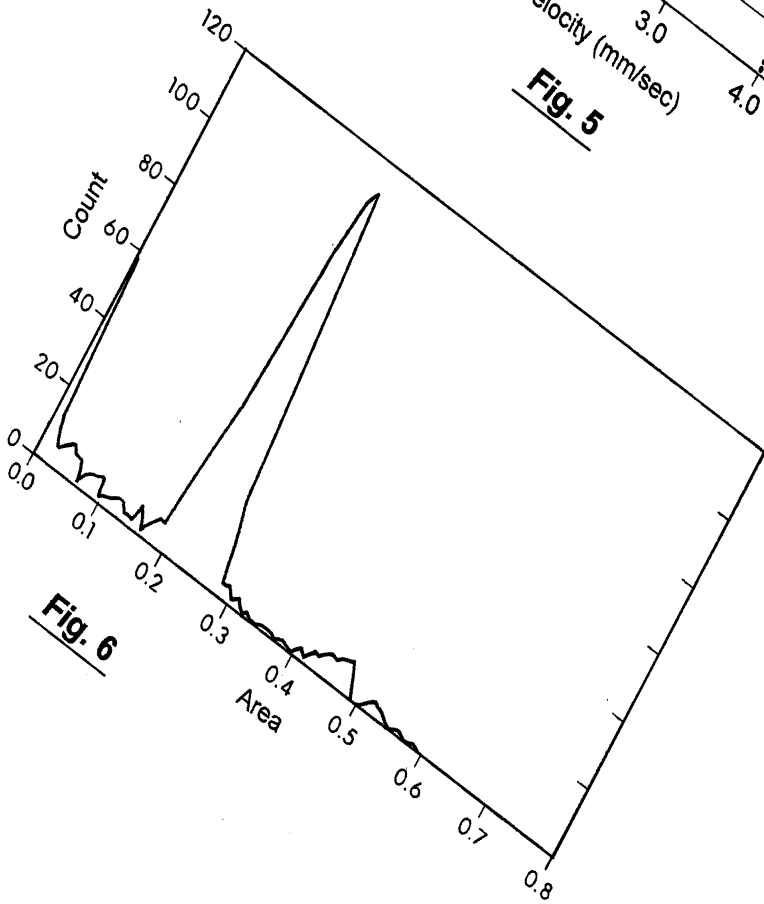


Fig. 6

ChDiv

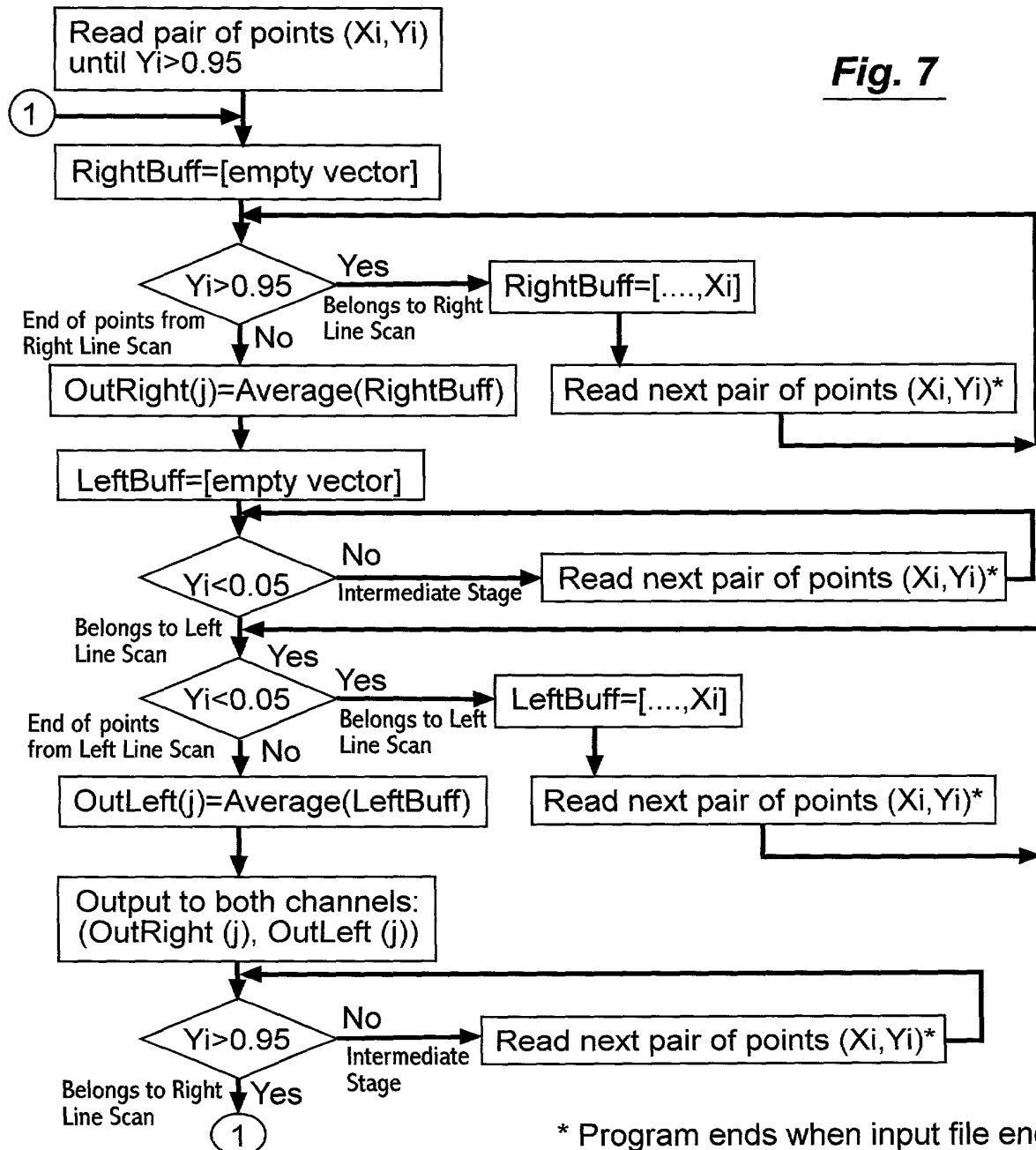
Input - two vectors: $Y(i)$ - channel 1 - square wave
- chopping signal, $0 \leq Y_i \leq 1$ $X(i)$ - channel 2 -
fluorescence raw data - from the detecting region
(both line scan)

Usually Sampled
at 40KHz

Output - two vectors: $OutRight(j)$ - fluorescence from
Right Line scan $OutLeft(j)$ - fluorescence from Left
Line scan

Usually Sampled
at 5KHz

*The sampling rate of the output channels allways equals the frequency of
the chopping signal*



ArV1Analyzer

Input: two files (one for each line scan).

Each file contain 2 vectors one of the Positions ($P(i)$) and the other has the corresponding Area ($A(i)$)

Output: three vectors - Area, TimeDiff (inversely proportional to velocity), Position

Position Parameters that can be determined - MinTimeDiff, Mas/timeDiff

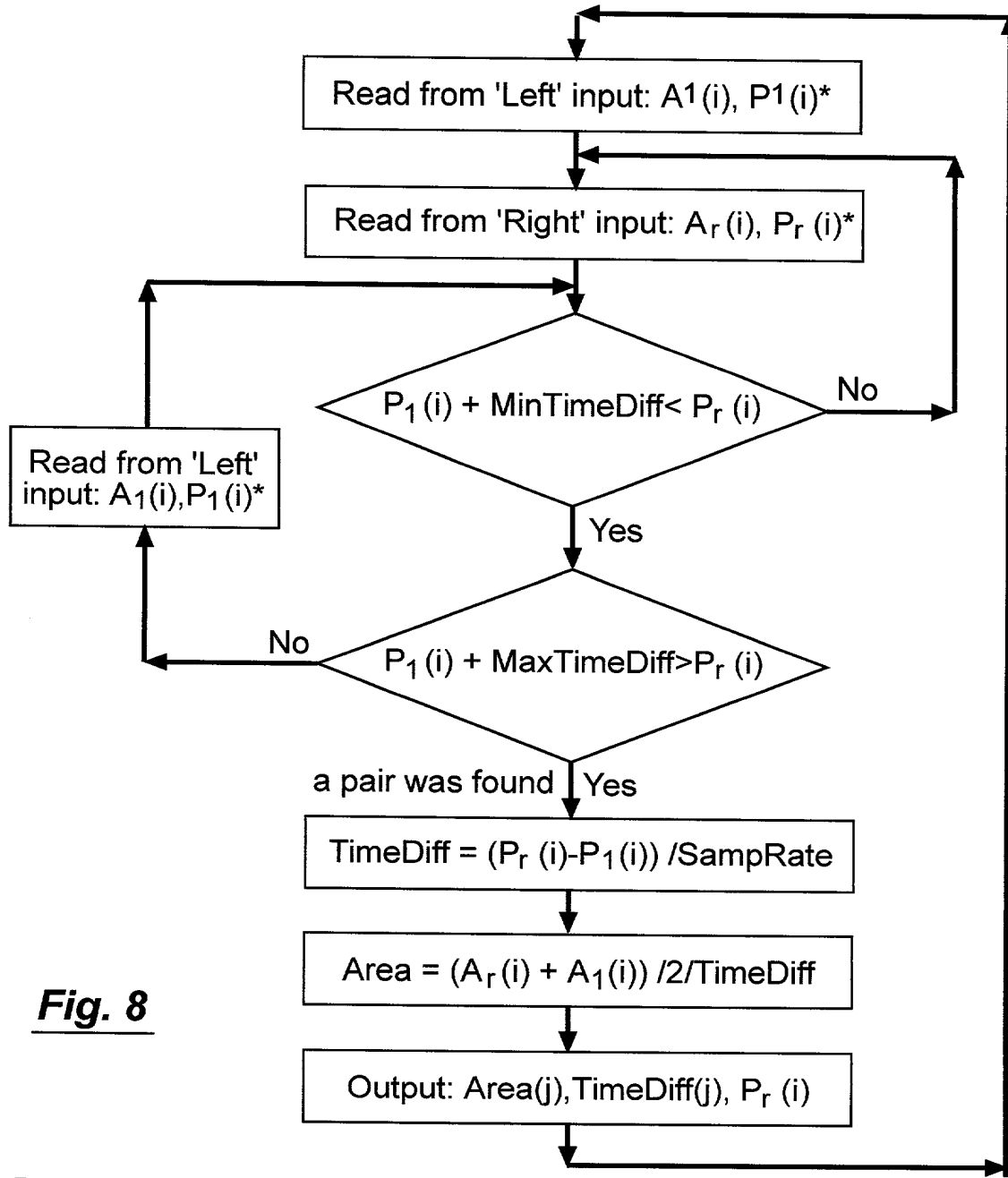


Fig. 8

Position is presented in point number and not time

TimeDiff is in Seconds and is inversely proportional to the velocity

*Program ends when one of the input files ends